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APPLICATION NO. FILING DATE FIRST NAMED INVENTOR ATTORNEY DOCKET NO.

09/095,385 06/10/98 M

MORRISON

30435.45USU1

HM12/0915

EXAMINER

GATES & COOPER HOWARD HUDHES CENTER 6701 CENTER DRIVE WEST, SUITE 1050 LOS ANGELES CA 90045 ZEMAN, M

ART UNIT

PAPER NUMBER

1643

10

DATE MAILED:

09/15/99

Please find below and/or attached an Office communication concerning this application or proceeding.

**Commissioner of Patents and Trademarks** 

	Application No.		Applicant(s)	
Office Action Summary	09/095,385		MORRISON ET AL.	
	Examiner		Art Unit	
	Mary K Zeman		1643	
The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply				
A SHORTENED STATUTORY PERIOD FOR REPLY THE MAILING DATE OF THIS COMMUNICATION.	/ IS SET TO EXF	PIRE <u>1</u> MONTH(S	S) FROM	
<ul> <li>Extensions of time may be available under the provisions of 37 after SIX (6) MONTHS from the mailing date of this communical.</li> <li>If the period for reply specified above is less than thirty (30) days be considered timely.</li> <li>If NO period for reply is specified above, the maximum statutory communication.</li> <li>Failure to reply within the set or extended period for reply will, by Status</li> </ul>	cation. s, a reply within the s period will apply and	tatutory minimum of I will expire SIX (6) M	thirty (30) days will	ailing date of this
1) Responsive to communication(s) filed on 10 S	September 1998 .			
2a) ☐ This action is <b>FINAL</b> . 2b) ☑ This action is non-final.				
3) Since this application is in condition for allowance except for formal matters, prosecution as to the ments is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.				
Disposition of Claims  4) □ Claim(s) 1-27 is/are pending in the application  4a) Of the above claim(s) is/are withdray  5) □ Claim(s) is/are allowed.  6) □ Claim(s) is/are rejected.  7) □ Claim(s) is/are objected to.  8) □ Claims 1-27 are subject to restriction and/or e  Application Papers  9) □ The specification is objected to by the Examine 10) □ The drawing(s) filed on is/are objected to 11) □ The proposed drawing correction filed on 12) □ The oath or declaration is objected to by the Examine 12) □ The oath or declaration is objected to by the Examine 12) □ The oath or declaration is objected to by the Examine 12) □ The oath or declaration is objected to by the Examine 12) □ The oath or declaration is objected to by the Examine 12) □ The oath or declaration is objected to by the Examine 12) □ The oath or declaration is objected to by the Examine 13.	wn from considerelection requirements.  er.  o by the Examine  _ is: a) ☐ approv	ent. er.	roved.	
Priority under 35 U.S.C. § 119				
13) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of the CERTIFI  1. received. 2. received in Application No. (Series Code 3. received in this National Stage applicatio * See the attached detailed Office action for a list of	IED copies of the  e / Serial Number  on from the Intern  of the certified co	priority docume  ) ational Bureau (I	nts have been: PCT Rule 17.2(a	a)).
14) Acknowledgement is made of a claim for domestic priority under 35 U.S.C. & 119(e).				
Attachment(s)				
<ul> <li>14) Notice of References Cited (PTO-892)</li> <li>15) Notice of Draftsperson's Patent Drawing Review (PTO-948)</li> <li>16) Information Disclosure Statement(s) (PTO-1449) Paper No(s)</li> </ul>	17)		/ (PTO-413) Paper I Patent Application (I	• •

Application/Control Number: 09/095,385

Art Unit: 1643

## Election/Restrictions

Restriction to one of the following inventions is required under 35 U.S.C. 121:

- Claims 1-13, drawn to recombinant methods of producing antibodies, classified in class 435, subclass 70.21.
- II. Claims 14 and 15, drawn to an antibody and a pharmaceutical composition, classified in class 424, subclass 147.1.
- III. Claims 16-21, drawn to methods of preventing infection, classified in class 424, subclass 204.1.
- IV. Claims 22-27, drawn to methods of treating an existing infection, classified in class 424, subclass 204.1.

The inventions are distinct, each from the other because of the following reasons:

Inventions I and II are related as process of making and product made. The inventions are distinct if either or both of the following can be shown: (1) that the process as claimed can be used to make other and materially different product or (2) that the product as claimed can be made by another and materially different process (MPEP § 806.05(f)). In the instant case secretory antibodies can be isolated from non-recombinant hybridomas expressing them.

Inventions II and (III/IV) are related as product and processes of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP)

Application/Control Number: 09/095,385

Art Unit: 1643

§ 806.05(h)). In the instant case the antibodies can be used in ELISA methods and for purification of the target polypeptide.

Inventions III and IV are related in that they use the same composition in the method however the methods have very different target populations and end results. Invention III would be given to a target population NOT infected with the relevant virus, to prevent infection, while Invention IV would be given to patients already infected with the virus, to treat the infection.

Invention I is patentably distinct from Inventions III and IV, as it is a separate and distinct method, having differing method steps, and resulting in differing outcomes. Invention I is drawn to producing antibodies *in vitro*, while Inventions III and IV are related to *in vivo* immunization processes.

Because these inventions are distinct for the reasons given above and have acquired a separate status in the art as shown by their different classification, restriction for examination purposes as indicated is proper.

Because these inventions are distinct for the reasons given above and the search required for Group I is not required for Group III and IV, restriction for examination purposes as indicated is proper.

This application contains claims directed to the following patentably distinct species of the claimed invention: If groups III or IV are selected Applicant must elect the pathogen from the following list: bacteria, virus, mycoplasma, yeast or parasite. If a virus is chosen as a species, a further election from the following is required: HIV, RSV, flu virus, or cold virus.

Application/Control Number: 09/095,385

Art Unit: 1643

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claims 16 and 22 are generic.

Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

Art Unit: 1643

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a petition under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

1. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Mary K Zeman whose telephone number is (703) 305-7133. The examiner can be reached between the hours of 7:30 am and 5:00 pm Monday through Thursday, and on alternate Fridays.

If attempts to reach the examiner by telephone are unsuccessful, Donna Wortman, Primary Examiner, can be reached on (703) 308-1032.

The fax number for this Art Unit is (703) 305-7401.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the Group receptionist whose telephone number is (703) 308-0196.

Mkz

September 14, 1999

DONNA WORTMAN PRIMARY EXAMINER